NEWS 42

Jan 29

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                  PCTFULL has been reloaded
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                  FOREGE no longer contains STANDARDS file segment
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                  USAN to be reloaded July 28, 2002;
NEWS 13 Jul 22
                   saved answer sets no longer valid
                   Enhanced polymer searching in REGISTRY
NEWS 14 Jul 29
                  NETFIRST to be removed from STN
NEWS 15 Jul 30
NEWS 16 Aug 08
                  CANCERLIT reload
                   PHARMAMarketLetter(PHARMAML) - new on STN
NEWS 17 Aug 08
                  NTIS has been reloaded and enhanced
NEWS 18 Aug 08
                  Aquatic Toxicity Information Retrieval (AQUIRE)
NEWS 19 Aug 19
                   now available on STN
                  IFIPAT, IFICDB, and IFIUDB have been reloaded
NEWS 20 Aug 19
                  The MEDLINE file segment of TOXCENTER has been reloaded
NEWS 21 Aug 19
                  Sequence searching in REGISTRY enhanced
NEWS 22 Aug 26
                   JAPIO has been reloaded and enhanced
NEWS 23 Sep 03
NEWS 24 Sep 16 Experimental properties added to the REGISTRY file
                  CA Section Thesaurus available in CAPLUS and CA
NEWS 25 Sep 16
                  CASREACT Enriched with Reactions from 1907 to 1985
NEWS 26 Oct 01
NEWS 27 Oct 21 EVENTLINE has been reloaded
NEWS 28 Oct 24 BEILSTEIN adds new search fields
NEWS 29 Oct 24 Nutraceuticals International (NUTRACEUT) now available on STN
NEWS 30 Oct 25 MEDLINE SDI run of October 8, 2002
NEWS 31 Nov 18 DKILIT has been renamed APOLLIT
NEWS 32 Nov 25 More calculated properties added to REGISTRY
                  TIBKAT will be removed from STN
NEWS 33 Dec 02
NEWS 34 Dec 04
                  CSA files on STN
                  PCTFULL now covers WP/PCT Applications from 1978 to date
NEWS 35 Dec 17
                  TOXCENTER enhanced with additional content
NEWS 36 Dec 17
                  Adis Clinical Trials Insight now available on STN
NEWS 37 Dec 17
NEWS 38 Dec 30
                  ISMEC no longer available
                  Indexing added to some pre-1967 records in CA/CAPLUS
NEWS 39 Jan 13
                  NUTRACEUT offering one free connect hour in February 2003
NEWS 40 Jan 21
                  PHARMAML offering one free connect hour in February 2003
NEWS 41
         Jan 21
                   Simultaneous left and right truncation added to COMPENDEX,
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118 L2 AND ACETONE

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L8 ANSWER 1 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 1
2001:89039 Document No. 134:212358 Dimension reduction and source identification for multispecies groundwater contamination. Duffy, C. J.; Brandes, D. (Department of Civil and Environmental Engineering, The Pennsylvania State University, University Park, PA, 16802, USA). Journal of Contaminant Hydrology, 48(1-2), 151-165 (English) 2001. CODEN: JCOHE6. ISSN: 0169-7722. Publisher: Elsevier Science B.V..

Assessment of chem. contamination at large industrial complexes with long AΒ and sometimes unknown histories of operation represents a challenging environmental problem. The spatial and temporal complexity of the contaminant may be due to changes in prodn. processes, differences in the chem. transport, and the phys. heterogeneity of the soil and aquifer materials. Traditional mapping techniques are of limited value for sites where dozens of chems. with diverse transport characteristics may be scattered over large spatial areas without documentation of disposal histories. In this context, a site with a long and largely undocumented disposal history of shallow groundwater contamination is examd. using principal component anal. (PCA). The dominant chem. groups and chem. modes at the site were identified. PCA results indicate that 5 primary and three transition chem. groups can be identified in the space of the 1st 3 eigenvectors of the correlation matrix, which account for 61% of the total variance of the data. These groups represent a significant redn. in the dimension of the original data (116 chems.). It is shown that each group represents a class of chems. with similar chemo-dynamic properties and/or environmental response. The groups are mapped back onto the site map to infer delineation of contaminant source areas for each class of compds. The approach serves as a preliminary step in subsurface characterization, and a data redn. strategy for source identification, subsurface modeling and remediation planning.

L8 ANSWER 2 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 2
2000:805407 Document No. 134:20790 Pollutant-Specific Scale of Multimedia
Models and Its Implications for the Potential Dose. Hertwich, Edgar G.;
McKone, Thomas E. (LCA-Laboratory, Norwegian University of Science and
Technology, Trondheim, 7491, Norway). Environmental Science and
Technology, 35(1), 142-148 (English) 2001. CODEN: ESTHAG. ISSN:
0013-936X. Publisher: American Chemical Society.

The spatial range is a generic indicator for how far pollutants are likely to travel. It also indicates the appropriate, pollutant-specific area of a multimedia model, which is the square of the spatial range. Formulations of the spatial range can be based on advective or dispersive transport. They differ in whether they take the extent and shape of the earth's surface into account. We suggest the common element of the different approaches is that all account for the persistence and mobility of pollutants. The mobility is the expected travel speed and depends on the partitioning. This paper extends the concept of a pollutant-specific

model scale through the introduction of a characteristic atm. scale height. It is the height of the atm. that would be needed to contain all the pollutant if the entire atm. had ground-level concn., taking into account deposition and degrdn. We define the spatial range as the expected advection-driven travel distance of a pollutant mol. released to a specific compartment. This novel anal. formulation is more comprehensive but encompasses all previous advection-based proposals of a spatial range. We evaluate the spatial range and scale height of 288 chems. for releases to air, surface water, and surface soil. We find a strong correlation between the spatial range for air releases and the scale height because both depend on persistence. We investigate the effect of the spatial scale on calcns. of the human toxicity potential, a screening-level risk indicator based on toxicity and potential dose. The product of model area and potential dose is found to be the same for calcns. using a fixed model area and those using the pollutant-specific spatial scale. The introduction of the scale height, however, can change the potential dose by more than 1 order of magnitude.

- L8 ANSWER 3 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 3
 1999:807646 Document No. 132:39919 Statistical comparison of leachate from hazardous, codisposal, and municipal solid waste landfills. Gibbons, Robert D.; Dolan, David G.; May, Helen; O'Leary, Kevin; O'Hara, Richard (University of Illinois at Chicago, IL, USA). Ground Water Monitoring and Remediation, 19(4), 57-72 (English) 1999. CODEN: GWMREV. ISSN: 1069-3629. Publisher: Ground Water Publishing.
- There has been considerable debate regarding the chem. characterization of AΒ landfill leachate in general and the comparison of various types of landfill leachate (e.g., hazardous, codisposal, and municipal) in particular. For example, the preamble to the US EPA Subtitle D regulation (40 CFR Parts 257 and 258) suggests that there are no significant differences between the no. and concn. of toxic constituents in hazardous vs. municipal solid waste landfill leachate. The purpose of this paper is to statistically test this hypothesis in a large leachate database comprising 1490 leachate samples from 283 sample points (i.e., monitoring location such as a leachate sump) in 93 landfill waste cells (i.e., a section of a facility that took a specific waste stream or collection of similar waste streams) from 48 sites with municipal, codisposal, or hazardous waste site histories. Results reveal clear differentiation between landfill leachate types, both in terms of constituents detected and their concns. The result is a classification function that can est. the probability that new leachate or groundwater sample was produced by the disposal of municipal, codisposal, or hazardous waste. This type of computation is illustrated, and applications of the model to Superfund cost-allocation problems are discussed.
- L8 ANSWER 4 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 4
 1998:666024 Document No. 129:299139 Toxicity tests in cell cultures for the purpose of predicting acute toxicity (LD50) and reducing the number of animal experiments. Halle, Willi (Forschungszentrum Juelich G.m.b.H., Juelich, D-52425, Germany). Schriften des Forschungszentrums Juelich, Lebenswissenschaften/Life Sciences, 1, 1-92 (German) 1998. CODEN: SFLSF9. ISSN: 1433-5549. Publisher: Forschungszentrum Juelich GmbH.
- An in vitro procedure for the redn. of animal expts. for toxicity tests of drugs or chems. is presented. Cytotoxicity data from in vitro cultivated mammalian cell lines were compared with acute toxicity data to predict the acute toxicity effects of xenobiotics in lab. animals. The procedure is based on a comparison of IC50 values (IC50x) with LD50 values using linear regression anal. An enlarged registry (RC) of cytotoxicity is presented contg. cytotoxicity data (IC50x) from non-selected chems. and drugs, the acute oral and i.v. LD50 values (LD50 p.o. and LD50 i.v.) from rats and mice, and the phys.-chem. characteristics of the chems. For the substances of the RC, sorted according to their IC50x-LD50 p.o. pairs, the

linear regression parameters were: r = 0.672, intercept a = 0.625, and slope b = 0.435. For the IC50x-LD50 i.v. pairs, the same parameters were: r = 0.768, a = -0.201, and b = 0.480. Approx. 73% of the p.o. values and 78% of the i.v. values are localized in the LD50 dosage range around the regression lines defined by an empirical factor FG.ltoreq.log 5. This percentage factor characterizes the dosage range of LD50 deviating from the regression line by the min. and max. residuals .ltoreq.0.699. The reliability of the predictive procedure was secured by using different biometrical methods and by comparisons of literature results with the data pool in the RC. The allocation of chems. into the 4 toxicity classes of acute oral toxicity defined by EU regulations (OECD Guide-line 423) resulted an accuracy of 85% in predicting the toxicity classes of the RC-substances in comparison to the toxicity classes of the corresponding NIOSH LD50 values. A comparison of RC-data with the Acute Toxic Class(ATC) method for the classification of chems. into toxicity classes resulted in a combined RC-ATC-procedure allowing the redn. of animal nos. for allocating chems. to the EU toxicity classes by 30%.

- L8 ANSWER 5 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 5
 1997:613701 Document No. 127:238509 Development of structure
 biodegradability relationships (SBRs) for estimating half-lives of organic
 contaminants in soil systems. Govind, R.; Lei, L.; Tabak, H. (Department
 of Chemical Engineering, University of Cincinnati, Cincinnati, OH,
 45221-0171, USA). NATO ASI Series, Series 2: Environment,
 23(Biodegradability Prediction), 115-138 (English) 1996. CODEN: NSSEF4.
 ISSN: 1431-7141. Publisher: Kluwer.
- Knowledge of half-lives or biodegrdn. rate consts. in soil is useful for AΒ estg. the natural attenuation rates of contaminants due to microbial transformations and to make decisions regarding treatment action or no treatment with isolation of the contaminated site to minimize exposure to animal and human life. Half-life is defined as the time required for 50% of the contaminant to be biodegraded. Soil treatment is time consuming and expensive, and often for large isolated contaminated sites, relying on natural attenuation may be the most cost-effective soln. A neural network is trained to est. the range of half-lives for org. contaminants in soil. Soil half-life data, obtained from the literature for 258 chems. is correlated with 14 mol. fragments or indicators using a back-propagation neural network with 14 input nodes, 12 nodes in the hidden layer and 2 output nodes. A cross-validation method was used to test the neural network. The converged neural network produced <50% relative error for >80% of the chems. in the training set. Using a classification scheme of fast (half-life 1-7 days), moderately fast (half-life 7-28 days), slow (half-life 28-180 days) and resistant (half-life 180-365 days), the neural network was able to correctly classify >95% of the 258 chems. in the database.
- L8 ANSWER 6 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 6
 1994:662443 Document No. 121:262443 French limiting values for occupational exposure to chemicals. Anon. (Fr.). Cahiers de Notes Documentaires, 153, 557-74 (French) 1993. CODEN: CNDIBJ. ISSN: 0007-9952.
- AB Limit values (suggested limiting values and max. permissible values) for occupational exposure to chems., including carcinogens, which have been published by the French Labor Ministry are presented in one table. This table is preceded by information on the following points: monitoring of workplace atmospheres (sampling and anal.; aerosols); permitted values (definitions and aims; additivity convention; elements and compds.; limiting occupational exposure values; carcinogens); mandatory values; and values recommended by the French National Health Insurance Fund (CNAM).
- L8 ANSWER 7 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 7
 1993:65829 Document No. 118:65829 Air contaminants. (Occupational Safety and Health Administration, U. S. Dep. Labor, Washington, DC, 20210, USA).

Federal Register, 57(114, Bk. 2), 26002-601 (English) 12 Jun 1992. CODEN: FEREAC. ISSN: 0097-6326.

Proposed amendments of existing air contaminant stds. for the maritime and AB construction industries and extension of air contaminant stds. to agricultural employees (only employees of farms with >10 nonfamily employees are covered) are given under the Federal Occupational Safety and Health Administration. Tables that indicated transitional limits, based on established threshold limit values, indication of skin protection needs, proposed time-weighted av. exposure (any 8-h work shift for 40-h week), short-term exposure limit (15-min time-weighted av.), ceiling (exposure during any part of the work day, or if instantaneous monitoring is not feasible, the 15-min time-weighted av.), and/or skin protection needs are given for the shipyard, marine terminal and longshoring, construction, and agricultural industries. Extensive data on health effects of the substances to be regulated and preliminary regulatory impact analyses are given for general industry and the specific industrial sectors.

L8 ANSWER 8 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 8
1990:83178 Document No. 112:83178 Reportable quantity adjustments; delisting of ammonium thiosulfate. (United States Environmental Protection Agency, Washington, DC, 20460, USA). Federal Register, 54(155), 33426-84 (English) 14 Aug 1989. CODEN: FEREAC. ISSN: 0097-6326.

Under the Federal Comprehensive Environmental Response, Compensation, and Liability Act, the EPA is promulgating final reportable quantities (RQ) for 258 hazardous substances and hazardous waste streams. NH4 thiosulfate is removed from the list of hazardous substances since the median lethal concn. is well above 500 mg/L for aquatic toxicity. Also included in this final rule is replacement of the registered trademark Gelthane with the generic name difocal, as several companies manuf. this substance.

L8 ANSWER 9 OF 9 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 9

1989:218230 Document No. 110:218230 Air contaminants. (United States Occupational Safety and Health Administration, Washington, DC, 20210, USA). Federal Register, 54(12, Bk. 2), 2332-983 (English) 19 Jan 1989. CODEN: FEREAC. ISSN: 0097-6326.

AB Under the Federal Occupational Safety and Health act, OSHA is amending existing air containment stds. and setting new permissible exposure limits for toxic substances commonly used in the workplace.

=> d 12

- L2 ANSWER 1 OF 9082 MEDLINE
- AN 2002642415 MEDLINE
- DN 22288963 PubMed ID: 12402082
- TI Physiological and cellular responses of the 2,4-D degrading bacterium, Burkholderia cepacia YK-2, to the phenoxyherbicides 2,4-D and 2,4,5-T.
- AU Cho Yun-Seok; Kahng Hyung-Yeel; Kim Chi-Kyung; Kukor Jerome J; Oh Kye-Heon
- CS Department of Life Science, Soonchunhyang University, P.O. Box 97, Asan, Chung-Nam, 336-600, Korea.
- NC P42-ES-04911 (NIEHS)
- SO CURRENT MICROBIOLOGY, (2002 Dec) 45 (6) 415-22. Journal code: 7808448. ISSN: 0343-8651.
- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- os genbank-af232906
- EM 200301
- ED Entered STN: 20021029

Last Updated on STN: 20030125

Entered Medline: 20030124

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               2,4,T
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           204 L9 AND EXTRACTION
=> s 110 and acetone
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=> d 112 1-4 cbib abs
L12 ANSWER 1 OF 4 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
97227754 EMBASE Document No.: 1997227754. GC/MS (SIM) determination of 14
     pesticides including cyhexatin and 2,4,5-T in nuts. Kawasaki M.; Fukuhara
     K.; Katsumura R.; Takasaka N.; Uchiyama S.. M. Kawasaki, Hatano Research
     Institute Food, Drug Safety Center, 729-5, Ochiai, Hadano, Kanagawa 257,
     Japan. Journal of the Food Hygienic Society of Japan 38/3 (161-169)
     1997.
     Refs: 12.
     ISSN: 0015-6426. CODEN: SKEZAP. Pub. Country: Japan. Language: Japanese.
     Summary Language: English; Japanese.
     GC/MS with selected ion monitoring ((GC/MS (SIM)) has been developed as a
AΒ
     screening method for the determination of 14 pesticides residues including
     cyhexatin and 2,4,5-T in nuts. The pesticides were extracted with
     acetone from nuts, and then partitioned between n-hexane and
     acetonitrile. Clean-up was conducted by solid-phase extraction
     (SPE) with a Bond Elut Florisil column. Most of the pesticides were eluted
     in a mixture of n-hexane and acetone (17: 3) from the column,
     and the eluate was examined by GC/MS (SIM). Cyhexatin and 2,4,5-T were
     analyzed after derivatization and SPE clean-up. Recoveries of 14
     pesticides from 6 kinds of nuts were mostly over 80%. This method is
     effective for multi-residue analysis of pesticides in nuts.
L12 ANSWER 2 OF 4 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.DUPLICATE 1
97170517 EMBASE Document No.: 1997170517. Comparison between methyl and
     pentafluorobenzyl esterification for analysis of 9 phenoxy acid herbicides
     in agricultural products. Hirahara Y.; Miyata M.; Kamakura K.; Watanabe
     Y.; Takeda H.; Maeda K.; Tonogai Y.. Y. Tonogai, Osaka Branch, National
     Inst. of Health Sciences, 1-1-43 Hoenzaka, Chuo-ku, Osaka 540, Japan.
     Japanese Journal of Toxicology and Environmental Health 43/2 (129-139)
     1997.
     Refs: 22.
     ISSN: 0013-273X. CODEN: JJTHEC. Pub. Country: Japan. Language: Japanese.
     Summary Language: English; Japanese.
     Ester types (2,4-dichlorophenoxyacetic acid (2,4-D) ethyl, 2,4-D butyl and
AΒ
     MCPA-thioethyl) and free acid types (2,4-D, 2,4,5,-
     trichlorophenoxyacetic acid (2,4,5,-T)
     2-methyl-4-chlorophenoxyacetic acid (MCPA), m- chloroperbenzoic acid
     (MCPB), mecoprop and dichlorprop) of phenoxy acid herbicides in
     agricultural products were simultaneously extracted from samples with
     acetone for vegetables/fruits, and with acetonitrile for
     cereals/beans, and then re-extracted with ethyl acetate. Three ester types
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of herbicides extracted from the samples were cleaned up by Sep-Pak.RTM.

Florisil, and then determined by ECD-GC and GC-MS (SIM). Recoveries of the ester types of herbicides determined by GC and GC-MS spiked at 1.25 ppm to asparagus, taro, brown rice and corn were 61.4 76.9% and 50.8%-63.9%, respectively. Six free acid types of herbicides were esterified by pentafluorobenzyl bromide or diazomethane and determined by GC-MS (SIM) without cleaning up by solid phase extraction. Recoveries of the free acid types of herbicides esterified by pentafluorobenzylation and methylation spiked at 1.25 ppm to the samples were 68.5 98.5% and 53.7-88.1%, respectively. It was clarified that the concentration of pentafluorobenzyl bromide in acetone should be 5-10% to have sufficient recoveries for the esterification. Methylation was superior to pentafluorobenzylation for the simplicity of manipulation, but was inferior for the sensitivity of GC and GC-MS.

- L12 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2003 ACS

 1993:253530 Document No. 118:253530 Analysis of chlorophenoxy acids and other acidic contaminants in food crops. Hajslova, J.; Tahtah, W. H.; Jehlickova, Z.; Kocourek, V.; Cuhra, P. (Dep. Food Chem. Anal., Prague Inst. Chem. Technol., Prague, 166 28, Czech.). Science of the Total Environment, 132(2-3), 259-74 (English) 1993. CODEN: STENDL. ISSN: 0048-9697.
- AB Several chlorophenoxy acids and chlorinated phenols were detd. by means of gas chromatog. in contaminated samples of cereals. Extn. of the plant matrix with an acetone/water mixt. followed by alk. hydrolysis was suitable for isolation of both free and conjugated residues. The use of pentafluorobenzyl bromide for volatilization of analytes, despite an enhanced ECD response, cannot be recommended for routine anal. Methylation with either methanol/sulfuric acid or methanol/BF3 reagent can substitute for diazomethane-based esterification procedure. Mass fragmentog. provided the highest selectivity of detection, and good sensitivity, 5 ppb. Even Me derivs. of monochlorinated analytes, which could not be measured by GC/ECD, could be quantitated at this level.
- L12 ANSWER 4 OF 4 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
 79015504 EMBASE Document No.: 1979015504. Evaluation of silica gel and XAD-4
 as adsorbents for herbicides in air. Grover R.; Kerr L.A.. Herbicide
 Behav. Environm. Sect., Res. Stat., Agric. Canada Regina, Saskatchewan S4P
 3A2, Canada. Journal of Environmental Science and Health Part B
 Pesticides, Food Contaminants, and Agricultural Wastes 13/3 (311-321)
 1978.
- CODEN: JPFCD2. Pub. Country: United States. Language: English.

 Silica gel and XAD-4 resin were shown to be efficient solid adsorbents for sampling vapors of n-butyl and iso-octyl esters of (2,4-dichlorophenoxy)acetic acid (2,4-D), n-butyl ester of (2,4,5-trichlorophenoxy)acetic acid (2,4,5-T), octanoate ester of 3,5-dibromo-4-hydroxybenzonitrile (bromoxynil), S-(2,3,3-trichloroallyl)diisopropylthiocarbamate (triallate), and alpha.,.alpha.,-trifluoro-2, 6-dinitro-N,N-dipropyl-p-toluidine (trifluralin) herbicides in ambient air. The absorbed herbicides were readily extracted by shaking the adsorbent with methanol or by Soxhlet extraction with a 1:1 mixture of acetone and n-hexane and subsequently analyzed with electron-capture GLC. The overall efficiencies of the two adsorbents, for both trapping and ease of extraction, were in the 85 to 100% range.